## II. Claim Amendments

- 1. (Presently amended) A multicomponent vaccine for ruminants comprising a safe and an immunogenically effective combination of a protective antigen component from a at least six clostridial organism organisms, a protective antigen component from a at least one non-clostridial organism and an adjuvant, wherein the vaccine is in a low dose volume of about 3 ml or less.
- 2. (Presently amended) A multicomponent vaccine comprising a safe and an immunogenically effective combination of protective antigen components from at least seven clostridial organisms, a protective antigen component from a at least one non-clostridial organism and an adjuvant, wherein the vaccine is in a low dose volume of about 3 ml or less.
- 3. (Presently amended) The vaccine according to Claim 1, wherein the clostridial organism is selected from the group consisting of Cl. chauvoei, Cl. septicum, Cl. novyi, Cl. perfringens type C, Cl perfringens type D, Cl. sordellii, Cl. haemolyticum and Cl. tetani.
- 4. (Presently amended) The vaccine according to Claim 1, wherein said non-clostridial organism is selected from

- the group consisting of a Gram negative bacteria bacterium, a Gram positive bacteria bacterium, a virus, a parasite and a rickettsia.
- 5. (Presently amended) The vaccine according to Claim 4, wherein the non-clostridial organism is at least one Gram negative organism is bacterium selected from the group consisting of H. somnus, M. bovis, P. haemolytica, P. multocida, E. coli, S. typhimurium, Leptospira spp. and C. foetus.
- 6. (Presently amended) The vaccine according to Claim 5, wherein the Gram negative organism bacterium is H. somnus.
- 7. (Presently amended) The vaccine according to Claim 5, wherein the Gram negative organism bacterium is M. bovis.
- 8. (Presently amended) The vaccine according to Claim 4, wherein the non-clostridial organism is at least one virus is selected from the group consisting of infections infectious bovine rhinotracheitis virus, bovine virus viral diarrhea virus, parainfluenza type 3 virus, bovine respiratory syncytial virus and a combination of at least two thereof.
- 9. (Presently amended) The vaccine according to Claim 4, wherein the non-clostridial organism is at least one

parasite is selected from the group consisting of Neospora spp., Tritrichimonas foetus and Cryptosporidium bovis.

- 10. (Cancelled)
- 11. (Presently amended) The vaccine according to Claim 1, wherein the adjuvant is selected from the group consisting of a polymer, a block co-polymer, an oil-in-water emulsion, a water-in-oil emulsion,

  Al (OH)<sub>3</sub>, AlPO<sub>4</sub>, an extract of a bacterial cell wall, an extract of a plant, a liposome, Quil A and a combination thereof a saponin and a combination of at least two thereof.

## Claims 12-14 Cancelled

- 15. (Presently amended) The vaccine according to Claim

  14 Claim 3, wherein the 6 clostridial organisms are
  selected from the group consisting of Cl. chauvoei,

  Cl. septicum, Cl.\_novyi, Cl. perfringens type C, Cl.
  perfringens type D, Cl. haemolyticum and Cl.
  sordellii.
- 16. (Cancelled)
- 17. (Presently amended) The vaccine according to Claim

  16 Claim 2, wherein the 7 clostridial organisms are selected from the group consisting of Cl. chauvoei,

- Cl. septicum, Cl. novyi, Cl. perfringens type C, Cl. perfringens type D, Cl. sordellii, Cl. haemolyticum, and Cl. tetani.
- according to Claim 1, for ruminants comprising a safe and immunogenically effective combination of a wherein the protective antigen component from 6 clostridial organisms which are from Cl. chauvoei, Cl. septicum, Cl novyi, Cl. perfringens type C, Cl. perfringens, type D, and Cl. sordellii; a and the protective antigen component from a non-clostridial organism which is from H. somnus and an adjuvant, wherein the vaccine is in a low dose volume.
- 19. (Presently amended) A multicomponent The vaccine for ruminants comprising a safe and immunogenically effective combination of an according to claim 2, wherein the protective antigen component from 7 clostridial organisms which are is from Cl. chauvoei, Cl. septicum, Cl novyi, Cl. perfringens type C, Cl. perfringens, type D, Cl. haemolyticum and Cl. sordellii; a and the protective antigen component from a non-clostridial organism which is from H. somnus and an adjuvant, wherein the vaccine is in a low dose volume.

- 20. (Cancelled)
- 21. (Cancelled)
- 22. (Presently amended) A The multicomponent vaccine for ruminants according to Claim 4, wherein at least one comprising a safe and immunogenically effective combination of protective antigen components from a clostridial organism; a protective antigen component is from a virus and an adjuvant, wherein the vaccine is in a dose size of 3.
- 23. (Presently amended) A The multicomponent vaccine for ruminants according to Claim 22, wherein the comprising a safe and immunogenically effective combination of protective antigen components from a plurality of clostridial organisms, a protective antigen component from comprises a plurality of viruses and an adjuvant, wherein the vaccine is in a dose size of 3.0 mL or less.
- 24. (Presently amended) The vaccine according to Claim 23, wherein the clostridial organism is organisms are selected from the group consisting of Cl. chauvoei, Cl. septicum, Cl\_novyi, Cl. perfringens type C, Cl. perfringens, type D, Cl. sordellii, Cl. haemolyticum, and Cl. tetani.

- 25. (Presently amended) The vaccine according to Claim 23, wherein the viruses are selected from the group consisting of infectious bovine rhinotracheitis, parainfluenza type 3 virus, bovine virus viral diarrhea virus and bovine respiratory syncytial virus.
- 26. (Presently amended) The vaccine according to Claim 23, wherein the adjuvant is selected from the group consisting of a polymer, a block co-polymer, an oil-in-water <a href="mailto:emulsion">emulsion</a>, a water-in-oil <a href="mailto:emulsion">emulsion</a>, an extract of a plant and a combination of at least two thereof.

## 27. (Cancelled)

- 28. (Presently amended) The vaccine according to Claim 2, wherein the non-clostridial organism is selected from the group consisting of a Gram negative bacteria bacterium, a Gram positive bacteria bacterium, a virus, a parasite and a rickettsia.
- 29. (Presently amended) The vaccine according to Claim 28, wherein the non-clostridial organism is a Gram negative bacterium and said Gram negative organism bacterium is selected from the group consisting of H. somnus, M. bovis, P.\_haemolytica, P. multocida, E. coli, S. typhimurium, Leptospira spp. and C. foetus.

- 30. (Presently amended) The vaccine according to Claim
  28, wherein the non-clostridial organism is a virus
  and the virus is selected from the group consisting of
  infectious bovine rhinotracheitis, parainfluenza type
  3 virus, bovine virus viral diarrhea virus and bovine
  respiratory syncytial virus.
- 31. (Presently amended) The vaccine according to Claim

  28, wherein the non-clostridial organism is a parasite

  and the parasite is selected from the group consisting

  of Neospora spp., Tritrichimonas foetus and

  Cryptosporidia spp..
- 32. (Cancelled)
- 28, wherein the adjuvant is selected from the group consisting of a polymer, a block polymer, an oil-in-water emulsion, a water-in-oil emulsion, an extract of a plant, a liposome and a combination of at least two thereof.

Claims 34 - 39 (Cancelled)

40. (Presently amended) A multicomponent The vaccine comprising a safe and immunogenically effective combination of a protective antigen component from according to claim 2, wherein the 7 clostridial

organisms which are Cl. chauvoei, Cl. septicum, Cl novyi, Cl. perfringens type C, Cl. perfringens type D, Cl. sordellii and Cl. haemolyticum; a and the protective antigen component from at least one non-clostridial organism is H. somnus or M. bovis and an adjuvant, wherein the vaccine is in a dose size of 3.0 mL or less.

- comprising a safe and immunogenically effective combination of a protective antigen component from 2 clostridial organisms which are selected from the group consisting of Cl. chauvoei, Cl. septicum, Cl novyi, Cl. perfringens type C, Cl. perfringens type D, Cl. sordellii, Cl. haemolyticum and Cl. tetani; a protective antigen component from viruses which are selected from the group consisting of infectious bovine rhinotracheitis virus, parainfluenza type 3 virus, bovine virus virus and an adjuvant, wherein the vaccine is in a dose size of 3.0 mL or less.
- 42. (Presently amended) A multicomponent vaccine comprising a safe and immunogenically effective combination of a protective antigen component from 6 clostridial organisms, which are Cl. chauvoei, Cl.

septicum, Cl novyi, Cl. perfringens type C, Cl.

perfringens type D and Cl. sordellii; a protective

antigen component from 4 viruses, which are infectious

bovine rhinotracheitis virus, parainfluenza type 3

virus, bovine virus viral diarrhea virus and bovine

respiratory syncytial virus, and an adjuvant, wherein

the vaccine is in a dose size of 3.0 mL or less.

## Claims 43-45 (Cancelled)

- 46. (New) A method of immunizing an animal comprising administering an effective amount of the vaccine of Claim 1.
- 47. (New) A method of immunizing an animal comprising administering an effective amount of the vaccine of Claim 2.